

Attorney Docket No.: 6544.200-US

**PATENT** 

## IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of: Persson et al.

Serial No.: 10/669,537

Group Art Unit: 1653

Filed: September 24, 2003

Examiner: To be assigned

For: Human Coagulation Factor VII Polypeptides

## **CERTIFICATE OF MAILING UNDER 37 CFR 1.8(a)**

Mail Stop Sequence Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Sir:

I hereby certify that the attached correspondence comprising:

- 1. Response to Notice to Comply with Sequence Rules
- 2. Sequence Listing
- 3. Copy of Notice to Comply with Sequence Rules
- 4. Disk Containing Sequence Listing

is being deposited with the United States Postal Service as first class mail in an envelope addressed to:

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## RESPONSE TO NOTICE COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES

MS: Sequence Commissioner for Patents P.O. Box 1450 Alexandria VA 22313-1450

Sir:

In response to the Notice to Comply With Requirements for Patent Applications Containing Nucleotide Sequence and/or Amino Acid Sequence Disclosure, a copy of which is enclosed, applicants enclose herewith the Sequence Listing for the above-captioned application and a 3.5" floppy disk containing the Sequence Listing. The content of the attached paper entitled "SEQUENCE LISTING" and of the accompanying identically labeled diskette is the same. Furthermore, the information contained in the attached "SEQUENCE LISTING" and the ASCII-encoded file is identical to the information in the specification as originally filed. No new matter is added.

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Serial No.; 10/669,537

No fee is due for this submission. However, please charge any fee, should it be required, to Novo Nordisk Pharmaceuticals, Inc., Deposit Account No. 14-1447. A duplicate of this sheet is enclosed.

Respectfully submitted,

Date: May 25, 2004

Reza Green, Reg. No. 38,475

Novo Nordisk Pharmaceuticals, Inc.

100 College Road West

Princeton, NJ 08540

(609) 987-5800

Use the following customer number for all correspondence regarding this application.

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**FORMALITIES LETTER** 

APPLICATION NUMBER

FILING OR 371 (c) DATE

FIRST NAMED APPLICANT

ATTORNEY DOCKET NUMBER

10/669.537

09/24/2003

Egon Persson

6544,200-US

**CONFIRMATION NO. 4625** 

Reza Green, Esq.

Novo Nordisk Pharmaceuticals, Inc.

100 College Road West Princeton, NJ 08540

DOCKET (check off

\*OC000000012192313\*

Date Mailed: 03/25/2004

NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE **DISCLOSURES** 

Filing Date Granted

Applicant is given TWO MONTHS FROM THE DATE OF THIS NOTICE within which to file the items indicated below to avoid abandonment. Extensions of time may be obtained under the provisions of 37 CFR 1.136(a).

• This application clearly fails to comply with the requirements of 37 CFR. 1.821-1.825. Applicant's attention is directed to the final rulemaking notice published at 55 FR 18230 (May 1, 1990), and 1114 OG 29 (May 15, 1990). If the effective filing date is on or after July 1, 1998, see the final rulemaking notice published at 63 FR 29620 (June 1, 1998) and 1211 OG 82 (June 23, 1998). If the effective filing date is on or after September 8, 2000, see the final rulemaking notice published in the Federal Register at 65 FR 54604 (September 8, 2000) and 1238 OG 145 (September 19, 2000). Applicant must provide an initial computer readable form (CRF) copy of the "Sequence Listing", an initial paper or compact disc copy of the "Sequence Listing", as well as an amendment directing its entry into the application. Applicant must also provide a statement that the content of the sequence listing information recorded in computer readable form is identical to the written (on paper or compact disc) sequence listing and, where applicable, includes no new matter, as required by 37 CFR 1.821(e), 1.821(f), 1.821(g), 1.825(b), or 1.825(d). If applicant desires the sequence listing in the instant application to be identical with that of another application on file in the U.S. Patent and Trademark Office, such request in accordance with 37 CFR 1.821(e) may be submitted in lieu of a new CRF.

For questions regarding compliance to these requirements, please contact:

MAR 29 2004

- For Rules Interpretation, call (703) 308-4216
- To Purchase Patentin Software, call (703) 306-2600
- For Patentin Software Program Help, call (703) 306-4119 or e-mail at patin21help@uspto.gov or patin3help@uspto.gov

Replies should be mailed to: Mail Stop Missing Parts

Commissioner for Patents

P.O. Box 1450

A copy of this notice <u>MUST</u> be returned with the reply.

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